10829139

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PASSWORD:
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NEWS 2		"Ask CAS" for self-help around the clock
NEWS 3	AUG 09	INSPEC enhanced with 1898-1968 archive
NEWS 4	AUG 28	ADISCTI Reloaded and Enhanced
NEWS 5	AUG 30	CA(SM)/CAplus(SM) Austrian patent law changes
NEWS 6	SEP 21	CA/CAplus fields enhanced with simultaneous left and right truncation
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NEWS EXPRESS NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.

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FILE 'HOME' ENTERED AT 18:18:00 ON 03 JAN 2007

=> fil reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

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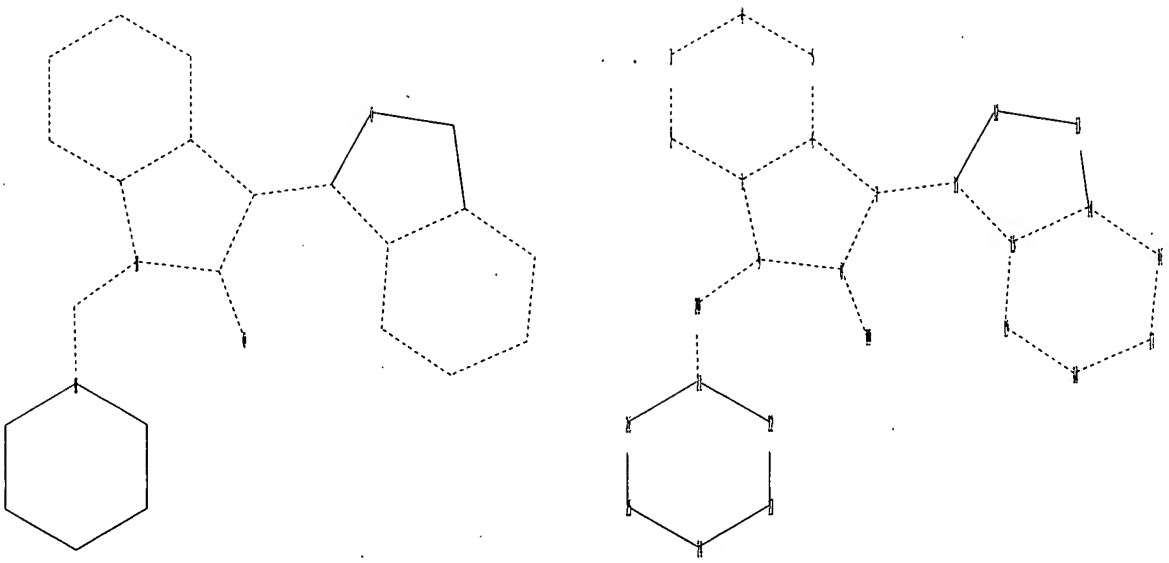
10829139



chain nodes :
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ring nodes :
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chain bonds :
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ring bonds :
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23-24 24-25 25-26

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom
20:CLASS 21:Atom 22:Atom 23:Atom 24:Atom 25:Atom 26:Atom

L1 STRUCTURE UPLOADED

=> d l1
L1 HAS NO ANSWERS
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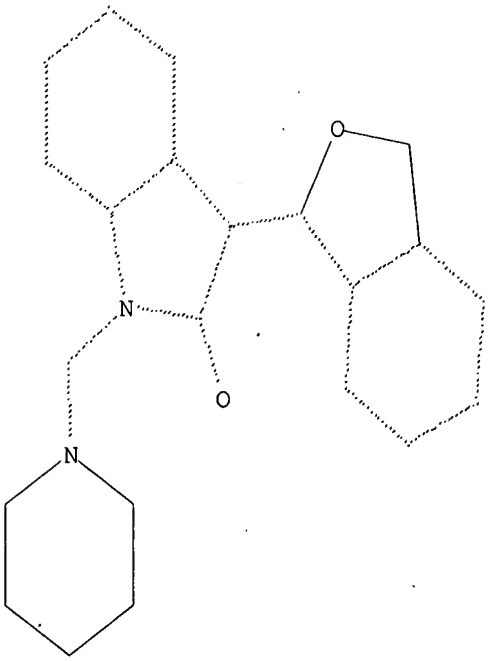
10829139



Structure attributes must be viewed using STN Express query preparation.

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100.0% PROCESSED       17 ITERATIONS                                0 ANSWERS
SEARCH TIME: 00.00.01

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                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:           93 TO      587
PROJECTED ANSWERS:               0 TO        0

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100.0% PROCESSED      322 ITERATIONS                                3 ANSWERS
SEARCH TIME: 00.00.01

L3            3 SEA SSS FUL L1

=> fil hcaplus
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                                                         ENTRY    SESSION
FULL ESTIMATED COST                                     172.55     172.76

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FILE COVERS 1907 - 3 Jan 2007 VOL 146 ISS 2
FILE LAST UPDATED: 2 Jan 2007 (20070102/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l3
L4 2 L3

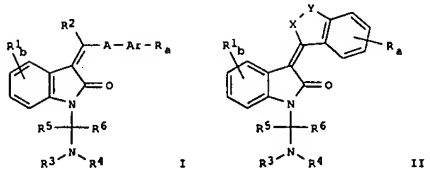
=> d ed ibib abs hitstr 1-2

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN
ED Entered STN: 28 May 2004
ACCESSION NUMBER: 2004:433773 .HCAPLUS
DOCUMENT NUMBER: 140:423586
TITLE: Preparation of dihydroindolones as tyrosine kinase inhibitors for the treatment of disease
INVENTOR(S): Andrews, Steven W.; Garst, Michael E.; Guo, Xialing; Hebert, Jonathan J.; Malone, Thomas; Wurster, Julie A.; Hull, Clarence Eugene
PATENT ASSIGNEE(S): Allergan, Inc., USA
SOURCE: U.S. Pat. Appl. Publ., 18 pp., Cont. of U.S. Ser. No. 306,975, abandoned.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

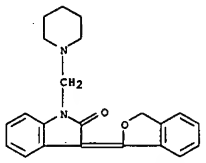
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004102509	A1	20040527	US 2003-389416	20030313
US 6747025	B2	20040608		
CA 2507780	A1	20040617	CA 2003-2507780	20031119
WO 2004050621	A2	20040617	WO 2003-US36988	20031119
WO 2004050621	A3	20040715		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
AU 2003295658	A1	20040623	AU 2003-295658	20031119
GB 2410744	A	20050810	GB 2005-11267	20031119
GB 2410744	B	20060412		
BR 2003016744	A	20051018	BR 2003-16744	20031119
JP 2006512400	T	20060413	JP 2004-570761	20031119
US 2004198802	A1	20041007	US 2004-829139	20040420
PRIORITY APPLN. INFO.:			US 2002-306975 B1	20021127
			US 2002-307097 A	20021127
			US 2003-389416 A	20030313
			WO 2003-US36988 W	20031119

OTHER SOURCE(S): MARPAT 140:423586
GI

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)

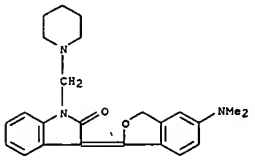


AB The title compds. [I or II; X = O, C(R2)2; Y = [C(R2)2]c; A = NR2, absent;
R1 = halo, OH, NO2, CN, etc.; R2 = H, alkyl, Ph, etc.; R = halo, (un)substituted hydrocarbyl; R3, R4 = H, (un)substituted hydrocarbyl; NR3R4 = (hetero)cyclic ring; R5, R6 = H, alkyl, aryl; a = 0-3; b = 0-3; c = 1-2] which are capable of modulating tyrosine kinase signal transduction
in order to regulate, modulate and/or inhibit abnormal cell proliferation,
were prepared Thus, reacting 4-morpholinoaniline with 3-(hydroxymethylene)-
1,3-dihydroindol-2-one (preparation given) in THF afforded 92% 3-[(4-morpholinophenylamino)-methylene]-1,3-dihydroindol-2-one which showed IC50 of 260 nM against VEGFR2 kinase.
IT 663903-86-4P 663903-90-0P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of dihydroindolones as tyrosine kinase inhibitors for treatment
of disease)
RN 663903-86-4 HCAPLUS
CN 2H-Indol-2-one, 1,3-dihydro-3-(1(3H)-isobenzofuranylidene)-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)

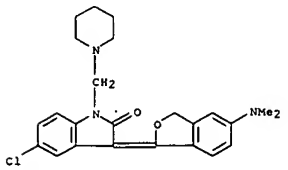


RN 663903-90-0 HCAPLUS

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)
CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of dihydroindolones as tyrosine kinase inhibitors for treatment
of disease)
RN 663903-91-1 HCAPLUS
CN 2H-Indol-2-one,
5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-
1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN
ED Entered STN: 04 Mar 2004
ACCESSION NUMBER: 2004:176559 HCAPLUS
DOCUMENT NUMBER: 140:210752
TITLE: Dihydroindolone compound tyrosine kinase inhibitors for the treatment of disease
INVENTOR(S): Andrews, Steven W.; Hebert, Jonathan J.
PATENT ASSIGNEE(S): Allergan, Inc., USA
SOURCE: U.S., 14 pp.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6699863	B1	20040302	US 2002-307097	20021127
CA 2507780	A1	20040617	CA 2003-2507780	20031119
WO 2004050621	A2	20040617	WO 2003-US36988	20031119
WO 2004050621	A3	20040715		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
AU 2003295658	A1	20040623	AU 2003-295658	20031119
GB 2410744	A	20050810	GB 2005-11267	20031119
GB 2410744	B	20060412		
BR 2003016744	A	20051018	BR 2003-16744	20031119
JP 2006512400	T	20060413	JP 2004-570761	20031119
PRIORITY APPLN. INFO.:			US 2002-306975 A	20021127
			US 2002-307097 A	20021127
			US 2003-389416 A	20030313
			WO 2003-US36988 W	20031119

OTHER SOURCE(S): MARPAT 140:210752
AB The invention discloses organic mols., especially dihydroindolone derivs. (preparation
described) capable of modulating tyrosine kinase signal transduction in order to regulate, modulate and/or inhibit abnormal cell proliferation.
IT 663903-86-4P 663903-90-0P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
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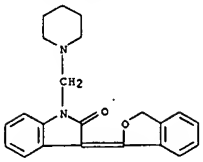
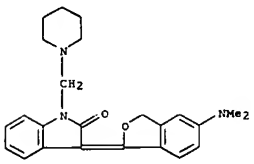
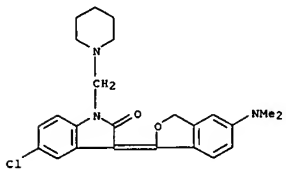


RN 663903-90-0 HCAPLUS
CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(dihydroindolone derivative tyrosine kinase inhibitors for treatment of disease)
RN 663903-91-1 HCAPLUS
CN 2H-Indol-2-one, 5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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=> d his

     (FILE 'HOME' ENTERED AT 18:18:00 ON 03 JAN 2007)

     FILE 'REGISTRY' ENTERED AT 18:18:11 ON 03 JAN 2007
L1              STRUCTURE UPLOADED
L2            0 S L1
L3            3 S L1 FULL

     FILE 'HCAPLUS' ENTERED AT 18:19:05 ON 03 JAN 2007
L4            2 S L3

=> log y
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY   SESSION
FULL ESTIMATED COST                                    15.74     188.50

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)          SINCE FILE      TOTAL
                                                       ENTRY    SESSION
CA SUBSCRIBER PRICE                                    -1.56      -1.56

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NEWS LOGIN Welcome Banner and News Items

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NEWS IPC8 For general information regarding STN implementation of IPC 8
NEWS X25 X.25 communication option no longer available

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FULL ESTIMATED COST	0.21	0.21

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DICTIONARY FILE UPDATES: 2 JAN 2007 HIGHEST RN 916646-22-5

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http://www.cas.org/ONLINE/UG/regprops.html

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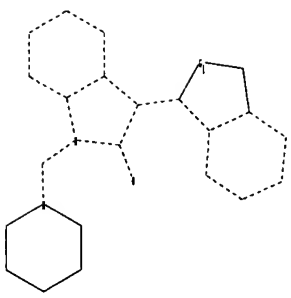
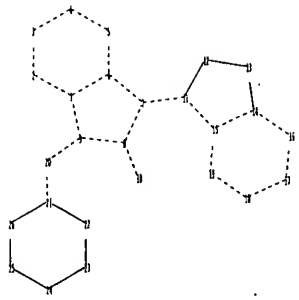
10829139s2





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ring nodes :
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23-24 24-25 25-26

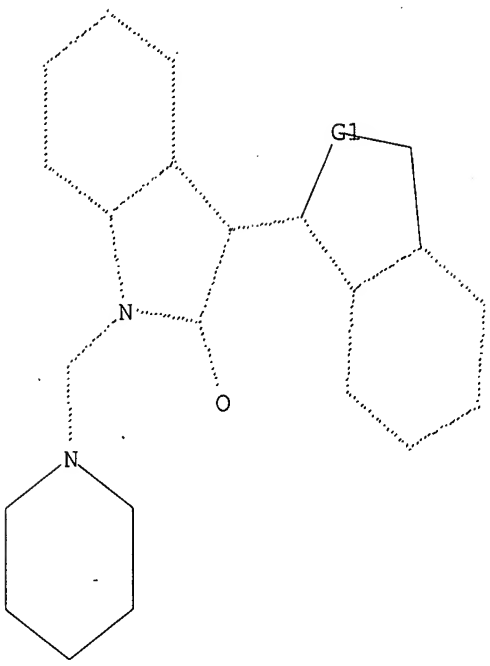
G1:C,O

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom
20:CLASS 21:Atom 22:Atom 23:Atom 24:Atom 25:Atom 26:Atom

L1 STRUCTURE UPLOADED

=> d l1
L1 HAS NO ANSWERS
L1 STR

10829139s2



G1 C,O

Structure attributes must be viewed using STN Express query preparation.

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SAMPLE SEARCH INITIATED 18:32:43 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -       36 TO ITERATE

100.0% PROCESSED       36 ITERATIONS                                0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:          360 TO     1080
PROJECTED ANSWERS:               0 TO        0

L2            0 SEA SSS SAM L1

=> s l1 full
FULL SEARCH INITIATED 18:32:49 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -      629 TO ITERATE

100.0% PROCESSED      629 ITERATIONS                                3 ANSWERS
SEARCH TIME: 00.00.01

L3            3 SEA SSS FUL L1

=> fil hcaplus
COST IN U.S. DOLLARS                          SINCE FILE      TOTAL
                                                 ENTRY    SESSION
FULL ESTIMATED COST                             172.10     172.31

FILE 'HCAPLUS' ENTERED AT 18:32:54 ON 03 JAN 2007
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
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10829139s2

FILE COVERS 1907 - 3 Jan 2007 VOL 146 ISS 2
FILE LAST UPDATED: 2 Jan 2007 (20070102/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l3
L4 2 L3

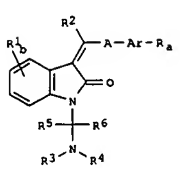
=> d ed ibib abs hitstr 1-2

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN
ED Entered STN: 28 May 2004
ACCESSION NUMBER: 2004:433773 HCAPLUS
DOCUMENT NUMBER: 140:423586
TITLE: Preparation of dihydroindolones as tyrosine kinase inhibitors for the treatment of disease
INVENTOR(S): Andrews, Steven W.; Garst, Michael E.; Guo, Xialing; Hebert, Jonathan J.; Malone, Thomas; Wurster, Julie A.; Hull, Clarence Eugene
PATENT ASSIGNEE(S): Allergan, Inc., USA
SOURCE: U.S. Pat. Appl. Publ., 18 pp., Cont. of U.S. Ser. No. 306,975, abandoned.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004102509	A1	20040527	US 2003-389416	20030313
US 6747025	B2	20040608		
CA 2507780	A1	20040617	CA 2003-2507780	20031119
WO 2004050621	A2	20040617	WO 2003-US36988	20031119
WO 2004050621	A3	20040715		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
AU 2003295658	A1	20040623	AU 2003-295658	20031119
GB 2410744	A	20050810	GB 2005-11267	20031119
GB 2410744	B	20060412		
BR 2003016744	A	20051018	BR 2003-16744	20031119
JP 2006512400	T	20060413	JP 2004-570761	20031119
US 2004198802	A1	20041007	US 2004-829139	20040420
PRIORITY APPLN. INFO.:			US 2002-306975	B1 20021127
			US 2002-307097	A 20021127
			US 2003-389416	A 20030313
			WO 2003-US36988	W 20031119

OTHER SOURCE(S): MARPAT 140:423586
GI

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)



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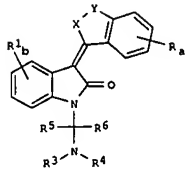
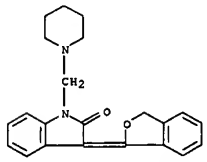


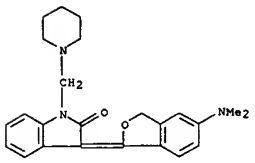
II

AB The title compds. [I or II; X = O, C(R2)2; Y = [C(R2)2]c; A = NR2, absent;
R1 = halo, OH, NO2, CN, etc.; R2 = H, alkyl, Ph, etc.; R = halo, (un)substituted hydrocarbyl; R3, R4 = H, (un)substituted hydrocarbyl; NR3R4 = (hetero)cyclic ring; R5, R6 = H, alkyl, aryl; a = 0-3; b = 0-3; c = 1-2] which are capable of modulating tyrosine kinase signal transduction
in order to regulate, modulate and/or inhibit abnormal cell proliferation,
were prepared Thus, reacting 4-morpholinoaniline with 3-(hydroxymethylene)-
1,3-dihydroindol-2-one (preparation given) in THF afforded 92% 3-[(4-morpholinophenylamino)-methylene]-1,3-dihydroindol-2-one which showed IC50 of 260 nM against VEGFR2 kinase.
IT 663903-86-4P 663903-90-0P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of dihydroindolones as tyrosine kinase inhibitors for treatment
of disease)
RN 663903-86-4 HCAPLUS
CN 2H-Indol-2-one, 1,3-dihydro-3-(1(3H)-isobenzofuranylidene)-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)

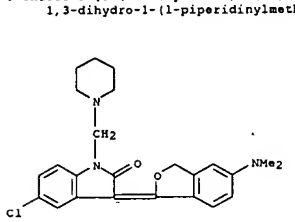


RN 663903-90-0 HCAPLUS

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)
CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of dihydroindolones as tyrosine kinase inhibitors for treatment
of disease)
RN 663903-91-1 HCAPLUS
CN 2H-Indol-2-one,
5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-
1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



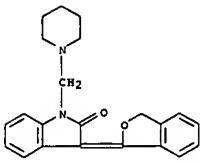
REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN
ED Entered STN: 04 Mar 2004
ACCESSION NUMBER: 2004:176559 HCAPLUS
DOCUMENT NUMBER: 140:210752
TITLE: Dihydroindolone compound tyrosine kinase inhibitors for the treatment of disease
INVENTOR(S): Andrews, Steven W.; Hebert, Jonathan J.
PATENT ASSIGNEE(S): Allergan, Inc., USA
SOURCE: U.S., 14 pp.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

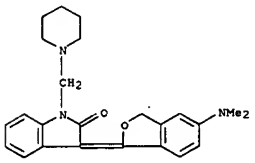
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6699863	B1	20040302	US 2002-307097	20021127
CA 2507780	A1	20040617	CA 2003-2507780	20031119
WO 2004050621	A2	20040617	WO 2003-US36988	20031119
WO 2004050621	A3	20040715		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
AU 2003295658	A1	20040623	AU 2003-295658	20031119
GB 2410744	A	20050810	GB 2005-11267	20031119
GB 2410744	B	20060412		
BR 2003016744	A	20051018	BR 2003-16744	20031119
JP 2006512400	T	20060413	JP 2004-570761	20031119
PRIORITY APPLN. INFO.:			US 2002-306975	A 20021127
			US 2002-307097	A 20021127
			US 2003-389416	A 20030313
			WO 2003-US36988	W 20031119

OTHER SOURCE(S): MARPAT 140:210752
AB The invention discloses organic mols., especially dihydroindolone derivs. (preparation
described) capable of modulating tyrosine kinase signal transduction in order to regulate, modulate and/or inhibit abnormal cell proliferation.
IT 663903-86-4P 663903-90-0P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(dihydroindolone derivative tyrosine kinase inhibitors for treatment of
disease)
RN 663903-86-4 HCAPLUS
CN 2H-Indol-2-one, 1,3-dihydro-3-(1(3H)-isobenzofuranylidene)-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)

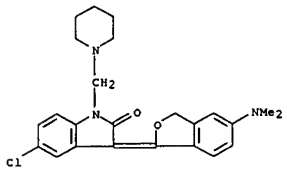
L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)



RN 663903-90-0 HCAPLUS
CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(dihydroindolone derivative tyrosine kinase inhibitors for treatment of disease)
RN 663903-91-1 HCAPLUS
CN 2H-Indol-2-one, 5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)

REFERENCE COUNT: 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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=> log y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	13.14	185.45

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-1.56	-1.56

STN INTERNATIONAL LOGOFF AT 18:33:17 ON 03 JAN 2007